

Appendix G.

Development of a Target Compound List (TCL)

G-1. Introduction

The two most common purposes for performing air monitoring at HTRW sites are to (1) provide information on HAPs concentrations for use in a site's overall hazard assessment and (2) assess the status of compliance with applicable Federal, state and local air pollution regulations. Both of these purposes can have a significant influence on the selection of sampling methods and the design of sampling programs.

As discussed earlier, CERCLA requires that a hazard assessment be performed at both Superfund and Corps HTRW sites, including an evaluation of the inhalation route of exposure. The usual approach for performing a hazard assessment is to make use of "risk assessment" techniques. The EPA's Superfund Public Health Evaluation Manual defines an acceptable level of cancer risk as being in the range of 10^{-7} to 10^{-4} . The EPA defines this as the acceptable level of risk for an adult exposed to maximum predicted ambient air concentration for a 70-year period, 24 hours per day. A 10^{-7} risk is a 1-in-10-million chance of death from cancer, whereas a 10^{-4} risk is a 1-in-10 thousand chance of death from cancer. Consequently, an FFMS at an HTRW site must be capable of measuring fence-line contaminant concentrations corresponding to risks within the 10^{-7} to 10^{-4} range.

In addition to the need for performing risk assessments, air sampling may also be required to determine the status of the HTRW site and its compliance with applicable regulations, defined in CERCLA as "applicable or relevant and appropriate" requirements (ARARs). An ARAR is a promulgated regulatory requirement at either the state or Federal levels of government (e.g., a National Ambient Air Quality Standard or a state air emission standard). ARARs apply to emissions from the HTRW site itself as well as to emissions from any remedial operations at the site.

In addressing the regulatory needs of the state agencies, EPA found a need to assist remediation programs in the identification of most probable analytes found at Superfund and HTRW sites. The objective of EPA developing a target compound list (TCL) was to help prioritize analytes of concern so applicable sampling and analytical methods could be identified and used in quantitating emissions to 10^{-6} risk levels.

Since no generally accepted list of HAPs existed, EPA developed a master list based upon the Hazardous Substances Priority Lists (HSPLs) and augmented with 60 additional HAPs selected from other authoritative lists (e.g., the Superfund Public Health Evaluation manual, the California Air Resources Board list of carcinogens, and lists published by the USEPA Office of Air Quality Planning and Standards).

After the master list was compiled, a simple scheme to rank these analytes in order of importance as HAPs at Superfund and HTRW sites was developed. The most important factors considered in developing this scheme were:

- Health effects of the analyte.

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- EPA, Corps, and state needs for regulating the analyte.
- Regulatory importance of the analyte.
- Potential for human exposure during site activities.
- Availability of sampling/analytical methods and reference standards for quantitating the analyte.

G-2. Health Effects

In considering health effects, a toxic compound list developed by EPA's Pollutant Assessment Branch (PAB) was used. This list is maintained within EPA's Office of Air Quality Planning and Standards (OAQPS). PAB also maintains a separate list of compound involving "cancer potency slopes" which in most cases are based upon the ingestion route of exposure. Because in many cases these cancer potency slopes have been, and will continue to be, converted to inhalation factors for use in HAPs risk assessments, these data were included in the assessment and ranking of health effects.

For noncarcinogens, lists maintained by EPA's noncarcinogen workgroup were used. These are compounds for which EPA has determined a need for the development of "reference dose" (RfD) values. RfD's are used by EPA as threshold values in evaluating noncarcinogenic health effect. For other compounds on the list which were not described by any of the above date, various health effects indicators such as threshold limit values, and as a last resort, reportable quantity date from SARA Title III, were relied upon.

G-3. EPA, Corps, and State Needs

In assessing EPA, Corps, and state needs for sampling guidance and analytical methods for specific HAPs, a questionnaire was developed and sent to interested parties to determine important HAPs of concern. The respondents provided lists of important HAPs, and the frequency with which specific compounds were of interest.

The response from the questionnaire was supplemented by a data base developed by the National Air Toxics Information Clearinghouse as an indicator of State regulatory activity for specific HAPs. For the various States regulating on the basis of acceptable ambient levels (AALs), the frequency of occurrence of regulations for specific chemicals was the third most important ranking criterion.

G-4. Regulatory Lists

Frequency of occurrence on lists of hazardous materials was also considered to be a useful ranking indicator. The California Air Resources Board (ARB) publishes a "Lists of Lists" which shows the frequency with which specific chemicals are listed in 12 authoritative lists of HAPs. The New York Air Guide II also categorizes specific air toxics compounds as high, medium, or low toxicity. SARA Title III, Section 313, also lists hazardous pollutants. Frequency of occurrence in each of these lists was used as an indicator of the relative importance of these compounds.

G-5. Potential for Human Exposure

Indicators for the potential for human exposure were incorporated by considering both the frequency of occurrence at Superfund and HTRW sites and the volatility of each of the listed compounds. Frequency of occurrence at Superfund sites was obtained directly from the August 1988 list entitled "Frequency Distribution of Substances Present at Final and Proposed NPL Sites." A volatility ranking number between 0.5 and 3 for each compound was derived from boiling point and/or vapor pressure data, as available. These indicators are generally considered to represent the potential for human exposure through the air pathway at Superfund sites.

G-6. Availability of Analytical Methods and Reference Standards

To complete the ranking process, each of the candidate chemicals on the expanded master list was entered into a Lotus 1-2-3 spreadsheet and arrayed with corresponding numerical data describing each of the 10 ranking criteria. A ranking index algorithm (RIA) was devised which would position the maximum value of each of the ranking criteria terms in its relative weighted position. The algorithm for ranking of the target compounds is:

$$\text{RIA} = 10\text{G} + 11.3\text{B} + 120\text{M} + 7.5\text{D} + 23.3\text{F} + 10\text{K} + 20\text{L} + 40\text{E} + 35\text{C} + 15\text{J}$$

Explanation of the development and derivation of term values can be found in Chapter 3.

As illustrated in Chapter 3, the RIA was designated as the sum of the descriptors terms. The complete target compound list developed for the Corps and EPA nationwide for Superfund sites utilizing the above algorithm consist of approximately 257 target compounds. Of the 257 compounds, 43 percent are volatiles thus having vapor pressure greater than 0.1 mm Hg. Approximately 32.4 percent of the target compound list are classified as semi-volatiles with vapor pressure ranging from 10^{-1} to 10^{-7} mmHg. Finally, metals comprise approximately 28 percent of the target compound list. The full target compound list of 257 compounds, marked in importance as determined by the RIA, is provided below in Table G-1.

Table G-1. EPA's Superfund Target Compound List (TCL)

Ranking (1)	Air Toxics (2)	CAA or 1990 (3)	CLASSI- FICATION Avail. (4)	STD's (5)	AVAILABLE SAMPLING/ANALYTICAL METHODS (6)	DEMONSTRATED SAMPLING METHOD DETECTION LIMITS (7)					UNIT RISK SLOPE CONCENTRATION AT 10 ⁻⁶ (8)	APPROXIMATE AIR RISK SPECIFIC CONCENTRATION AT 10 ⁻⁶ (9)	REFERENCE AMBIENT LEVELS (RAIS) (ug/m ³)/ppb (10)
						Sep-PAK (a)	Canister (b)	Tenax (c)	Filter (d)	PUF/XA D ² (e)			
(Abbreviation explanation at end of table)													
1	*Vinyl Chloride	+	V	+ N	C-GCMS		0.42	(a)			4.2x10 ⁻⁵	0.23/0.00898	0.0460/0.0180
2	*Trichloroethylene	+	V	+ N	TIC-GCMS		0.47	0.08			1.3x10 ⁻⁵	7890/1431	1.5380/0.2852
3	*Cadmium	+	M	NNT	FICAP, F-XR				0		1.8x10 ⁻³	.00555/0.0001196	0.0012/0.0003
4	*Chloroform	+	V	+ N	TIC-GCMS		0.37	0.23			2.3x10 ⁻⁵	0.43/0.008806	0.0860/0.0176
5	*Benzene	+	V	+ N	TIC-GCMS		0.53	2.9			8.3x10 ⁻⁶	.120/0.03756	0.2400/0.0751
6	*Carbon Tetrachloride	+	V	+ N,F7	TIC-GCMS		0.41	0.17			1.5x10 ⁻⁵	0.67/0.010648	0.1340/0.0213
7	*Arsenic	+	M	NNT	FICAP, F-XR				0		4.3x10 ⁻³	.00023/0.000751	
8	*Tetrachloroethylene	+	V	+ N	TIC-GCMS		0.57	2.6			9.5x10 ⁻⁷	1.05/0.15479	2.1000/0.3096
9	*Chromium	+	M	NNT	FICAP				0.002		1.2x10 ⁻²	.000093/0.000039	0.0002/0.0001
10	*Mercury	+	M	NNT	FICAP, CV-AA, ACM				0.008				0.0118/0.0014
11	*Beryllium	+	M	NNT	FICAP, F-XR				0		2.4x10 ⁻³	.0004/0.00108	0.0020/0.0054
12	*Selenium	+	M	NNT	FICAP, F-XR		(b)	(a)	0.001				0.5400/0.1872
13	*Nickel (Subsulfide)	+	M	N	FICAP, F-XR				0.002		4.8x10 ⁻⁴	0.0021	0.0060/0.0025
14	*Heptachlorocycloheptachloroperoxide	+	P	NNT	PUF-GC/ED, PUF-GCMS					0.01	1.3x10 ⁻³	.00077/0.000504	0.0020/0.0001
15	1,1-Dichloroethane	+	V	+ N	TIC-GCMS						5.0x10 ⁻⁵	.02/0.005044	0.0400/0.0101
16	*Acrylonitrile	+	V	+ F7	TIC-GCMS		0.1	(a)			6.8x10 ⁻⁵	0.0150/0.00812	0.0300/0.0138
17	*Benzo(a)pyrene		SV	NNT	PUF/XAD ² -HPLC, PUF/XAD ² -GC/MS, SPAD ² -HPLC/UV					0	1.7x10 ⁻³	.00058/0.000562	1.14/0.1105
18	*1,2-Dichloroethane	+	V	+ N	TIC-GCMS		0.39	3.8			2.6x10 ⁻⁵	0.038/0.0093877	0.0760/0.0188
19	Chlorobenzene	+	V	+ N	TIC-GCMS		0.48	1.7					10.02/1.722
20	Lead	+	M	NNT	FICAP, F-XR				0.0187				0.14/0.0185
21	*Formaldehyde	+	V	N	SEP-IC, ACM	0.012	(b)	(c)			6.1x10 ⁻⁶	184/0.13553	0.328
22	1,1,1-Trichloroethane	+	V	+ N	TIC-GCMS		0.42	1.7					0.0046/0.0007
23	*1,1,2-Trichloroethane	+	V	G,N,F7	TIC-GCMS		0.38	2.1			1.6x10 ⁻⁵	.062/0.01136	0.124/0.0227
24	*Chlordane	+	P	NNT	PUF-GC/ED, PUF-GCMS		(b)	(a)		0.01	3.7x10 ⁻⁴	.0027/0.00018	0.0054/0.0003
25	*1,1,2,2-Tetrachloroethane	+	V	G, F7	TIC-GCMS		0.66	6.5			5.8x10 ⁻⁵	.017/0.002476	0.034/0.0005
26	Barium		M	NNT	FICAP, F-XR				0				0.0200/0.0036
27	Ethylbenzene	+	V	+ N	TIC-GCMS		0.44	1.6					0.0152/0.0035
28	*PCBs	+	P	NNT	PUF/XAD ² -GC/MS, PUF/XAD ² -HRGC/HRMS		(b)	(a)		1.0x10 ⁻⁶	1.2x10 ⁻³	0.00089	0.001
29	*Asbestos	+	V	N	F-MICR								
30	*Toxaphene	+	P	N	PUF-GC/ED, PUF-GCMS						2.4x10 ⁻¹	0.000004	
31	*Methylene chloride	+	V	+ N	TIC-GCMS		(b)	(a)		0.01	3.2x10 ⁻⁴	.0031/0.0001832	1.447/0.0855
32	Manganese	+	M	NNT	FICAP, F-XR		0.73	(a)			4.1x10 ⁻⁶	2.44/0.70235	4.8800/1.4047
33	1,2,4-Trichlorobenzene	+	V	N, F7	TIC-GCMS		(b)	(c)	0				16.60007/3878
34	*Styrene	+	V	+ N	TIC-GCMS		0.45	0.13			2.8x10 ⁻⁷	3.48/0.81703	18.00007/2.4253
35	1,1-Dichloroethane	+	V	+ N	TIC-GCMS		0.51	5.7					1000.00007/247.04

Table G-1. (Continued)

Ranking (1)	Air Toxics (2)	CAA CLASSI- FICATION (3)	STD's Avail (4)	AVAILABLE SAMPLING/ANALYTICAL METHODS (5)	DEMONSTRATED SAMPLING METHODOLOGY (S) AND APPROACHABLE METHOD DETECTION LIMITS (7)				UNIT RISK SLOPE (6) (g/m ³) ⁻¹	APPROXIMATE AIR RISK SPECIFIC CONCENTRATION AT 10 ⁻⁵ (8)	REFERENCE AMBIENT LEVELS (RA) (s) (g/m ³) (ppb)	
					Sep-PAK (d) ug/m ³	Canister (c) ppb	Tenax (e) ng	Filter (f) ug/m ³				PUF-XA DT ² (h) ug/m ³
(Abbreviation explanation at end of table)												
37	Naphthalene	+	SV	N, G, NT	PUFXAD ² -HPLC, PUFXAD ² -GC/MS, XAD ² -HPLC/UV		(b)	(c)	0		14.25702/7169	
38	*Ethylene oxide	+	V	N	CT-GC/EC, CT-GC/FID		(b)	(a)	1.0x10 ⁻⁴	0.010/0.0557	0.02	
39	Toluene	+	V	+N	TIC-GCMS		0.4	2			10.24602/7169	
40	Xylenes (o, m and p)	+	V	+N, F7	TIC-GCMS		0.31	0.5			11.81002/72	
41	1,2-Dichloropropane	+	V	+N	TIC-GCMS		0.34	4				
42	1,2-Dichlorobenzene	+	V	G, +F7	TIC-GCMS		0.32	12.4			81.8330/13.8101	
43	1,2-Dibromothane	+	V	+	TIC-GCMS			3.3			2.462/0.3204	
44	*1,3-Butadiene	+	V	+	TIC-GCMS		0.66	(a)		4.8x10 ⁻⁷	4.2000/1.8985	
45	Thallium	+	M	N, NT	F-ICAP, F-GFAA, F-XR				0.0209	2.176/98089	0.5700/0.0854	
46	Zinc	+	M	N, NT	F-ICAP, F-AA, F-XR				0.002		18.008/7314	
47	Copper	+	M	N, NT	F-ICAP, F-AA, F-XR				0		5.002/1930	
48	*Propylene oxide	+	V	N	CT-GC/FID, CT-GCMS			(a)		1.2x10 ⁻⁴	1.88	
49	Acetone	+	V	+N, F7	TIC-GCMS, SEP-IC	0.0237	3.3	(c)			180.087/3554	
50	Chloroethane	+	V	G, +N, F7	C-GC/EC, TIC-GCMS, CT-GCMS		0.37	(c)			718.3590/272.1	
51	Phenol	+	SV	N	IMP-HPLC		(c)	(c)			20.015	
52	3,3-Dichlorobenzidine	+	SV	N	FISG-HPLC/UV		(b)	(a)			0.197	
53	*2,3,7,8-tetrachlorodibenzo-p-dioxin	+	SV	N	PUFXAD ² -HRGC/HRMS		(b)	(a)	1x10 ⁻⁶	0.280/0.019745	0.06	
54	2-Butanone	+	V	+N	TIC-GCMS			(c)				
55	*Nitrobenzene	+	SV	N	TIC-GCMS, PUF-XAD ² -GC/MS		(c)	(c)		8.33/1.854382	4.0000/0.7944	
56	*Dieldrin/Aldrin	+	P	N, NT	PUF-GC/EC, PUF-GCMS		(b)	(a)	0.01	0.0021/0.000141	0.0004	
57	Hexachlorocyclopentadiene	+	SV	N	PUF-GC/EC, PUF-GCMS			(a)	0		0.1200/0.0108	
58	Acrolein	+	V	N	TIC-GCMS, SEP-IC	0.0229	(b)	(a)			0.3000/0.1308	
59	*Hexachlorobenzene	+	SV	N, NT	PUF-GC/EC, PUF-GCMS		(b)	(a)	0	0.022/0.0001717	0.0040/0.0003	
60	Antimony	+	M	N, NT	F-ICAP, F-AA, F-XR				0.002		1.3390/0.2889	
61	*Benzyl chloride	+	V	N, +F7	TIC-GCMS		(c)	(c)		1.2x10 ⁻⁵	0.8330/0.16031	
62	*Pentachlorophenol	+	SV	N	PUF-GC/EC, PUF-GCMS		(b)	(a)	0.173	3.9x10 ⁻⁷	5.0000/0.4590	
63	Carbon Disulfide	+	V	N	C-GCMS, GB-GCMS, ACM		(c)	(a)			0.270/0.0887	
64	*4,4-DDE DDT, DDD	+	P	N, NT	PUFXAD ² -GC/EC, PUFXAD ² -GC/MS		(b)	(a)	0.002	0.1030/0.00079	2.0600/0.1421	
65	Hydrogen fluoride	+	V	N, G	DENUDER-IC, SILICA GEL-HPLC/UV ACM		(b)	(a)			0.679	
66	4-Methyl-2-pentanone	+	V	N, +F7	SEP-IC, TIC-GCMS		(c)	(c)			53.9100/13.16	
67	Cobalt	+	M	N, NT	F-ICAP, F-XR		(c)		0.004		0.570/2385	
68	Nickel carbonyl	+	V		FICT-GFAA			(a)				
69	Cis-1,3-Dichloropropene	+	V	G, N	TIC-GCMS	0.53		(c)				
70	Phosgene	+	V		IMP-HPLC, ACM			(a)			0.485	

Table G-1. (Continued)													
Ranking (1)	Air Toxics (2)	CAA of 1990 (3)	CLASSI- FICATION Avail. (4)	STDs Avail. (4)	AVAILABLE SAMPLING/ANALYTICAL METHODS (5)	DEMONSTRATED SAMPLING METHODOLOGY (5) AND APPROACHABLE METHOD DETECTION LIMITS (7)					UNIT RISK SLOPE ($\mu\text{g}/\text{m}^3$) (6)	APPROXIMATE AIR RISK SPECIFIC CONCENTRATION AT 10^{-6} (8)	REFERENCE LEVELS (RAIS) ($\mu\text{g}/\text{m}^3$)/ppb (9)
						Sec-PAK (a) $\mu\text{g}/\text{m}^3$	Canister (b) ppb	Terax (c) ng	Filter (d) $\mu\text{g}/\text{m}^3$	PUFA D 25 (e) $\mu\text{g}/\text{m}^3$			
(Abbreviation explanation at end of table)													
72	Nickel		M	NNT	FICAP, F-AA, F-XR								
73	Hydrazine	+	V		SFIRBK-HPLC/UV								
74	Fluorene		SV	NNT	PUF/XAD 25 -HPLC, PUF/XAD 25 -GC/MS, SXAD 25 -HPLC/UV		(b)	(a)		0			
75	1,4-Dichlorobenzene	+	V	G	TIC-GC/MS		0.12/0.79	0.5				0.36100/0600	
76	1,4-Dioxane	+	V	+	T-GC/MS		0.9	3.9				0.48000/1332	
77	Nickel sulfide		M	N	IMP-COL								
78	Ammonia		V	N/G	DENUDER-IC, ACM		(b)	(a)				4.736	
79	*Epichlorohydrin	+	V	N	T-GC/MS		(c)	(c)		2.7×10^{-6}	3700.097768		
80	*Acetaldehyde	+	V	N	IMP-HPLC/UV, SEP-IC	0.02				2.2×10^{-6}	0.450/246773		
81	Aniline	+	SV	N	IMP-COL, PUF-GC/MS		(c)	(c)					
82	*Hexachlorobutadiene	+	V	G	C-GC/MS		(c)	(c)		2.2×10^{-5}	0.045	2.9	
83	Methyl isocyanate	+	V		SXAD 25 -HPLC/UV		(b)	(a)					
84	Toluene diisocyanate	+	SV	N	GB-GC/FID		(b)	(a)					
85	Silver		M	NNT	FICAP, F-AA, F-XR				0			0.59800/0.1928	
86	Bromomethane	+	V	+N	TIC-GC/MS, CT-GC/MS		0.84	7.8				80.0000/15.4502	
87	Ethylamine		V		IMP-HPLC		(c)	(a)					
88	Trans 1,3-Dichloropropene	+	V	G/N	TIC-GC/MS		0.4	(c)					
89	Methoxychlor	+	P	N	PUF-GC/EC, PUF-GC/MS		(b)	(a)		0.01		56.97204/0300	
90	Dichlorodifluoromethane		V	+	C-GC/MS, CT-GC/MS		(c)	(c)				0.03000/0.0061	
91	Parathion	+	P	N	PUF-GC/MS, PUF-GC/EC, PUF-GC/NPD		(b)	(a)		0		0.570/0.0478	
92	Hydrogen sulfide		V	G	IMP-COL, GB-GC/FPD, ACM		(c)	(a)				3.79	
93	*Chloromethane	+	V	G+F7	C-GC/MS, CT-GC/MS		0.39	(a)		1.8×10^{-6}	0.555/0.28378	11.1000/5.3752	
94	*N-Nitrosodimethylamine	+	V	N	ADS-GC/MS, STN-GC/EC		(c)	(c)		5.4×10^{-3}	0.000190/0.000594		
95	Benzo(b)fluoranthene		SV	NNT	PUF/XAD 25 -HPLC, PUF/XAD 25 -GC/MS, PUF-GC/MS, PUF-GC/FID, SXAD 25 -HPLC/UV		(b)	(a)		0			
96	Radon	+	V	N	CT-RAD								
97	Cis-1,2-Dichloroethylene		V	G/N	TIC-GC/MS		0.060/25	(c)					
98	Fluoridesulfonate/HF		V	N/G	DENUDER-IC, SILICA GEL/HC, ACM							6.807	
99	Sulfur Dioxide		V	G	IMP-COL, GB-GC/FPD, ACM		(c)	(a)				28.5002/7568	
100	Methanol	+	V	G	TIC-GC/MS, TIC-GC/FID		20	(a)					
101	Bromodichloromethane		V	G+F7	TIC-GC/MS		0.48	(c)					
102	Hydrogen Arsenide		V	N	CT-GFAA								
103	Trisbromomethane	+	V	N+F7	TIC-GC/MS		0.48	(a)				28.5	
104	Acetonitrile	+	V	+	TIC-GC/MS		(c)	(c)				100.0000/69.5615	
105	1,3-Dichlorobenzene		V	G	TIC-GC/MS		0.070/44	(a)					

Table G-1. (Continued)														
Ranking (1)	Air Toxics (2)	CAA of (3)	CLASSI- FICATION (3)	STD's Avail. (4)	AVAILABLE SAMPLING/ANALYTICAL METHODS (6)	DEMONSTRATED SAMPLING METHODOLOGY (5) AND APPROACHABLE METHOD DETECTION LIMITS (7)					UNIT RISK SLOPE (UC/m ³) (8)	APPROXIMATE AIR RISK SPECIFIC CONCENTRATION AT 10 ⁶ (9)	REFERENCE AMBIENT LEVELS (RA)LS (ug/m ³ /ppb) (9)	
						Sep-PAK (a)	Canister (d)	Tenax (e)	Filter (f)	PUF/XA D ² (f)				
														ug/m ³ (a)
(Abbreviation explanation at end of table)														
107	Benzo(a)anthracene		SV	N/NT	PUF/XAD ² -HPLC, PUF/XAD ² -GC/MS, SXAD ² -HPLC/UV		(b)	(a)		0				
108	Pyridine		V	+	T-GC/MS, SXAD ² -GC/MS		(c)	(c)					4.000/1.2364	
109	BHC		P	N/NT	PUF-GC/EC, PUF-GC/MS					0.01				
110	Endosulfon		P	N	PUF-GC/EC, PUF-GC/MS		(b)	(a)					0.5690/0.0342	
111	Methoxyethanamine		V											
112	*3-chloro-1-propene	+	V	N	TIC-GC/MS, CT-GC/MS		(c)	(c)			5.5x10 ⁻⁶	13.18	363.6000/118.1639	
113	Dibenz(a,h)anthracene		SV	N/NT	PUF/XAD ² -HPLC, PUF-GC/MS, PUF-GC/EC, SXAD ² -HPLC/UV					0				
114	Boron		M	N/NT	F-ICAP, F-XR				0.004					
115	Benzo(k)fluoranthene		SV	N/NT	PUF/XAD ² -GC/MS, PUF-GC/EC, SXAD ² -HPLC/UV									
116	Dibromochloromethane		V	G+N/7	TIC-GC/MS			(c)					0.57	
117	Endrin aldehyde/endrin		P	N/NT	PUF-GC/EC, PUF-GC/MS					0.01				
118	Methyl Methacrylate	+	V	N	T-GC/MS, C-GC/MS									
119	Anthracene		SV	N/NT	PUF/XAD ² -HPLC, PUF-GC/MS, PUF-GC/EC, SXAD ² -HPLC/UV					0				
120	Mirex		P	N/NT	T-GC/MS, PUF-GC/EC, PUF-GC/MS					0.01				
121	Dibromochloropropane		V	N	CT-GC/MS									
122	Tetrahydrofuran		V	N	T-GC/MS, PUF-GC/MS, PUF-HRGC/HRMS, C-GC/MS			1.2					160.478	
123	Bromoethane		V	N	TIC-GC/MS		(b)	(c)						
124	2-chloro-1,3-Butadiene	+	V	G	TIC-GC/MS		0.38						0.890/3.132	
125	Vinyl Acetate	+	V	N	C-GC/MS		(c)	(c)					38.3060/10.88	
126	Sulfuric Acid		SV	N	DENUDER-C, B-HPLC, ACM								2.728	
127	4-Chloroaniline		SV	N	PUF-GC/MS									
128	DiChloromethylether		V											
129	Thorium		M	N	F-ICAP, F-XR				0.008					
130	*Trans-1,4-Dichlorobutene		V		TIC-GC/MS		(c)	(c)			2.8x10 ⁻³	0.00038		
131	Bromochloromethane		V	G	TIC-GC/MS		0.67	2.1					30.165	
132	*Benzidine	+	SV	N	F-HPLC/UV, PUF-GC/MS						6.7x10 ⁻²	.000015/0.00002		
133	Methacrylonitrile		V	N	T-GC/MS, C-GC/MS									
134	Propylene		V	G	T-GC/EC, TIC-GC/MS		0.29							
135	1,1,2-Trichloro-1,2,2-trifluoroethane		V	+	TIC-GC/MS									
136	Acenaphthylene		SV	NT	PUF/XAD ² -GC/MS					0.003				
137	Benzo(g,h,i)Perylene		SV	N	PUF-GC/EC, PUF/XAD ² -HPLC, PUF/XAD ² -GC/MS, SXAD ² -HPLC/UV		(b)	(c)		0				

Table G-1. (Continued)												
Ranking (1)	Air Toxics (2)	CAA of 1990 (3)	CLASSI- FICATION (3)	STD's Avail. (4)	AVAILABLE SAMPLING/ANALYTICAL METHODS (6)	DEMONSTRATED SAMPLING METHODOLOGY (S) AND APPROACHABLE METHOD DETECTION LIMITS (7)					APPROXIMATE AIR RISK SPECIFIC CONCENTRATION AT 10 ⁻⁵ (8)	REFERENCE AMBIENT LEVELS (RAIS) (ugm ³ /ppb) (9)
						Sep-PAK (a)	Canister (d)	Tenax (e)	Filter (f)	PUF/XA D ² (g)		
(Abbreviation explanation at end of table)												
139	Hydrogen Cyanide	+	V	N	IMP-COL, IMP-IC, ACM					ugm ³ /ppb	86	
140	Aldicarb		P	N	PUF-HPLC, PUF-GCMS, PUF-GCECD, SEP-IC, SEP-HPLC					0.003	4	
141	Furfural		V	N	T-GCMS, PUF-GCMS							
142	Phenanthrene		SV	N	PUF/XAD ² -GCMS, PUF/XAD ² -HPLC, PUF-GCECD, SXAD ² -HPLC/UV					0		
143	1,1-Dimethylhydrazine		V									
144	Zinc Oxide		M	N	F-ICAP, F-XR				0.001			
145	Polybrominated biphenyls		P	N	PUF/XAD ² -GCMS, PUF-GCECD					1x10 ⁻⁶		
146	Pyrene		SV	N	PUF/XAD ² -HPLC, PUF/XAD ² -GCMS, PUF-GCECD, SXAD ² -HPLC/UV		(c)	(c)		0		
147	Trichlorofluoromethane		V	+	T-GCMS, CT-GCMS		(c)					
148	1,2,3-Trichloropropane	+	V	N	T-GCMS		(c)	4.7				
149	1,2-Diphenylhydrazine	+	SV		F/IMP-HPLC/UV							
150	Uranium		M	N	F-ICAP							
151	*2,4,6-Trichlorophenol	+	SV	N	PUF-GCECD, PUF-GCMS, PUF-GCEC D					0.01	5.7x10 ⁻⁵	
152	2,4-Dinitrotoluene	+	SV	N	T-GCMS, PUF-GCMS							
153	2,4-Dichlorophenol		SV	N	T-GCECD, T-GCMS							
154	Isopropylbenzene	+	V	G	T-GCMS							
155	Methylene bis(phenyl isocyanate)		SV	N								
156	Indeno(1,2,3-cd)pyrene		SV	N	PUF/XAD ² -HPLC, PUF/XAD ² -GCMS, PUF-GCECD, SXAD ² -HPLC/UV					0		
157	Tin		M	N	F-ICAP, F-XR				0.03			
158	Molybdenum		M	N	F-ICAP, F-XR				0			
159	Dibenzofuran	+	SV	N	PUF/XAD ² -GCMS, PUF/XAD ² -HPLC, PUF-GCECD					1x10 ⁻⁶		
160	Cresols	+	SV	N	IMP-HPLC						24.061	
161	Chrysene		SV	N	PUF/XAD ² -HPLC, PUF/XAD ² -GCMS, PUF-GCECD, SXAD ² -HPLC/UV					0		
162	2-Methoxyethanol		V	N	C-GCMD							
163	Heptane		V	G	T-GCMS		0.176/78					
164	Acetic Anhydride		V		IMP-COL							
165	Malathion		P	N	PUF-GCMS, PUF-GCECD, PUF-GC/MS, PUF-GC/MS					0.01		
166	*Hexachloroethane	+	SV	N	T-GCECD, PUF-GCMS, SXAD ² -GCMS						4.0x10 ⁻⁵	
167	2,4,5-Trichlorophenol	+	SV	N	PUF/XAD ² -GCCECD, PUF/XAD ² -GCMS, PUF-HPLC					0.07	0.25/0.0258	

Table G-1. (Continued)												
Ranking (1)	Air Toxics (2)	CAA of 1990 (3)	STD's Avail (4)	AVAILABLE SAMPLING/ANALYTICAL METHODS (5)	DEMONSTRATED SAMPLING METHODOLOGY (S) AND APPROACHABLE METHOD DETECTION LIMITS (T)					UNIT RISK SLOPE (U)(m ⁻³) (6)	APPROXIMATE AIR RISK SPECIFIC CONCENTRATION AT 10 ⁻⁶ (8)	REFERENCE AMBIENT LEVELS (RALS) (ug/m ³)ppb (9)
					Ssp-PAK (g) ugm ³	Canister (d) ppb	Tenax (e) ng	Filter (f) ugm ³	PUF/XA D ⁺ (g) ugm ³			
(Abbreviation explanation at end of table)												
169	Diethyl nitrosamines	V	N	STN-GC/NPD,STN-GC/HECD								
170	Diethylphthalate	SV	N	PUF-GC/MS,SXAD ² -GC/MS								
171	Maleic anhydride	SV	N	SXAD ² -HPLC/UV								
172	2-Chlorophenol	V	N	S/GILICA GEL-HPLC/UV								
173	2-Chloropropane	V	N	TIC-GC/MS			3.4					
174	Strontium	M	N	F-ICAP, F-XR				0.002				
175	Ethylene diamine	V	N	STN-GC/NPD								
176	Chlorodibenzodioxins	SV	N	PUF-HRGC/HRMS								
177	B-Naphthylamine	SV	N	IMP-HPLC,PUF-GC/MS							1x10 ⁻⁸	
178	Bis(2-Ethylhexyl)phthalate	SV	N	PUF-GC/MS,SXAD ² -GC/MS								
179	2-Chloroethyl vinyl ether	V	N	T-GC/MS								
180	2-Ethoxyethanol	V	N	CT-GC/FID								
181	n-Nitrosodiphenylamine	SV	N	STN-GC/NPD								
182	Octane	V	G	TIC-GC/MS		0.05/0.28						
183	Chlorodifluoromethane	V	G	C-GC/MS,CT-GC/MS								
184	Isophorone	SV	N	T-GC/MS, PUF-GC/MS,SXAD ² -GC/MS								
185	2,4-D Salts & esters	SV	N	F-HPLC/UV								
186	*Di(n-octyl)phthalate	SV	N	PUF-GC/MS,SXAD ² -GC/MS							7.69/0.48148	
187	Nitrophenol	SV	N	PUF-GC/MS,SXAD ² -GC/MS								
188	Acenaphthene	SV	N	PUF/XAD ² -HPLC, PUF/XAD ² -GC/MS, PUF-GC/FID					0			
189	Bis(2-chloroethyl)ether	SV	N	SXAD ² -GC/MS,PUF-GC/MS								
190	Bromobenzene	V	N	T-GC/MS			14.1					
191	Benzoic acid	SV	N	SXAD ² -GC/MS								
192	Butylbenzylphthalate	SV	N	SXAD ² -GC/MS,PUF-GC/MS								
193	Fluoranthene	SV	N	PUF/XAD ² -HPLC, PUF/XAD ² -GC/MS, PUF-FID, SXAD ² -HPLC/UV					0			
194	Disulfoton	P	N	PUF-GC/HECD, PUF-HPLC, PUF-GC/NPD, PUF-GC/FID, PUF-GC/MS					0.01			
195	Methyl styrene	V		TIC-GC/MS								
196	2-Nitrophenol	SV	N	PUF-GC/HECD,PUF-GC/MS,PUF-HPLC, SXAD ² -GC/MS					0.01			
197	2,4-Dimethyl phenol	SV	N	SXAD ² -GC/MS,PUF-GC/MS								
198	Plutonium	M										
199	Benzaldehyde	V	N	SEP-IC,B-HPLC/UV	0.0433		5.9					
200	Dicyclopentadiene	SV	N	CT-GC/FID,SXAD ² -GC/MS								

Table G-1. (Continued)

Table G-1. (Continued)													
Ranking (1)	Air Toxics (2)	CAA of 1990	CLASSI- FICATION (3)	STD's Avail. (4)	AVAILABLE SAMPLING/ANALYTICAL METHODS (5)	DEMONSTRATED SAMPLING METHODOLOGY (6) AND APPROACHABLE METHOD DETECTION LIMITS (7)					UNIT RISK SLOPE ($\mu\text{g}/\text{m}^3$) (8)	APPROXIMATE AIR RISK SPECIFIC CONCENTRATION AT 10 ⁻³ (9)	REFERENCE AMBIENT LEVELS (RAAs) ($\mu\text{g}/\text{m}^3$)/ppb (10)
						Sep-PAK ($\mu\text{g}/\text{m}^3$)	Canister (g)	Tenax (g)	Filter (g)	PUFVA D-2 ($\mu\text{g}/\text{m}^3$)			
(Abbreviation explanation at end of table)													
203	2,6-Dinitrotoluene		SV	N	FIMP-HPLC/UV, SXAD ² -GC/MS								
204	Benzonitrile		SV	N	T-GC/MS, SXAD ² -GC/MS			1.3					
205	Ethylene glycol	+	SV	N	FICT-GC/FID								
206	2,4-Dinitrophenol	+	SV	N	SXAD ² -GC/MS, PUF-GC/MS								
207	n-Nitroso-n-propylamine		SV	N	STIN-GC/HECD								
208	1,2-Dimethylhydrazine		V										
209	Radium		M	N	F-ICAP Radiochemical Method								
210	Tritium		V		F-ICAP, F-XR								
211	2-Nitroaniline		SV	N	SXAD ² -GC/MS, PUF-GC/MS								
212	3-Nitroaniline		SV	N	SXAD ² -GC/MS, PUF-GC/MS								
213	Coke oven emissions	+	SV										
214	n-Pentane		V	G	TIC-GC/MS		0.280.85						
215	2-Hexanone		V	N	SEP-IC-T-GC/MS, B-HPLC/UV								
216	4-Nitrodiphenyl	+	SV	N	IMP-HPLC/UV, SXAD ² -GC/MS, PUF-GC/MS								
217	p-Biphenylamine		SV	N	SXAD ² -GC/MS, PUF-GC/MS								
218	Carbaryl	+	P	N	F-HPLC/UV								
219	2,4,5-trichlorophenoxyacetic acid	+	P	N	F-HPLC/UV								
220	Auramine		P		F-HPLC/UV								
221	*Coal Tars		SV		PUF-HPLC/UV								
222	Di-(n-butyl)phthalate		SV	N	SXAD ² -GC/MS, PUF-GC/MS					1x10 ⁻⁸	0.0016		
223	Cresote		SV		IMP-HPLC, PUF-HPLC, PUF-GC/MS, PUF-GC/FID					0.001			
224	Dimethylformamide	+	V	N	SILICA GEL-GC/FID								
225	Atrazine		P	N	F-HPLC/UV, PUF-HPLC/UV								
226	Acetophenone	+	SV	N	T-GC/MS, SEP-IC, PUF, SXAD ² -HPLC					0			
227	Di-(2-Chloroethoxy)methane		SV	N	PUF-GC/MS, SXAD ² -GC/MS								
228	Ethylene glycol monobutyl ether		V										
229	Cyclopentadiene		V		SICROMO 104-GC/FID								
230	Nitrates/nitrites		SV	N	DENUDER-IC								
231	4,6-dinitro-2-methylphenol		SV	N	SXAD ² -GC/MS, PUF-GC/MS								
232	Hexane	+	V	G	TIC-GC/MS		0.060.23						
233	Cyclohexanone		V	N	SEP-IC-T-GC/MS, B-HPLC/UV								
234	Cyclohexylamine		V	N	SILICA GEL-GC/NPD								
235	Dimethylphthalate	+	SV	N	SXAD ² -GC/MS, PUF-GC/MS								
236	Acetone Cyanohydrin		SV	N	SIPORAPAK-QS, GC/NPD								

Table G-1. (Continued)												
Ranking (1)	Air Toxics (2)	CAA of 1990 (3)	CLASSI- FICATION (4)	STD's Avail (5)	AVAILABLE SAMPLING/ANALYTICAL METHODS (6)	DEMONSTRATED SAMPLING METHODOLOGY (6) AND APPROACHABLE METHOD DETECTION LIMITS (7)				UNIT RISK SLOPE (Ug/m ³) (8)	APPROXIMATE AIR RISK SPECIFIC CONCENTRATION AT 10 ⁶ (9)	REFERENCE AMBIENT LEVELS (RA-LS) (ug/m ³)/ppb (10)
						Sep-PAK (a)	Canister (b)	Filter (c)	PUFA D ² (d)			
(Abbreviation explanation at end of table)												
238	1-bromobutane		V		T-GCMS	ug/m ³	ppb	ng	ug/m ³	ug/m ³	ug/m ³ /ppb	
239	1-bromo-4-phenoxybenzene		V									
240	4-chloro-3-methylphenol	+	SV	N	B-COL, SXAD ² -GC/MS							
241	2,4,5-TP		P	N	IMP-GC/ECD							
242	Phthalic anhydride	+	SV	N	F-HPLC/UV							
243	Pentachlorobenzene		P	N	FIS-GC/ECD,PUFAD ² -GC/MS							
244	4,4-Methylene-bis- (2-chloroaniline)	+	SV	N								
245	Propylene glycol monomethyl ether		V	N								
246	4-Methyl phenol		SV	N	SXAD ² -GC/MS,PUF-GCMS							
247	2-Methyl phenol		SV	N	SXAD ² -GC/MS,PUF-GCMS							
248	Benzyl alcohol		SV	N	SXAD ² -GC/MS,PUF-GCMS							
249	2-Methylnaphthalene		SV	N	T-GCMS, SXAD ² -GC/MS, PUF-GCMS				0			
250	*Nitrosomorpholine	+	V	N	STN-GC/ECD					2.5x10 ⁻⁵	0.0400.008423	
251	2,4,6-Trinitrobenzene		SV	N	F-GC/MS,SXAD ² -GC/ECD							
252	2,4,6-Trinitrophenylmethyl- inilamine		SV	N								
253	cyclohexane		SV		F-HPLC/UV							
254	Nitrosobenzene		SV									
255	Ethylene cyanohydrin		SV									
256	Propylene glycol		SV	N								
257	1,3,5-Trinitrobenzene		SV	N	SXAD ² -GC/MS							
(a) Not Amenable to Tenax Analysis												
(b) Not Amenable to Canister Analysis												
(c) No Detection Limits Available, but Feasible												
1 Reference to 40CFR 60.130												
2 Based on 10L Samples												

Abbreviations for Table G-1, EPA's Superfund Target Compound List

(1) As determined by EPA's RIA, discussed in Chapter 3.

(2) Those toxics that have unit risk numbers developed by U.S. Environmental Protection Agency and other agencies are indicated by an asterisk.

(3) Classification

V = Volatile air toxic compounds having vapor pressure above 10^{-1} mm Hg at standard conditions (20°C and 760 mm Hg).

SV = Volatile air toxic compounds having vapor pressure between 10^{-1} and 10^{-7} mm Hg at standard conditions (20°C and 760 mm Hg).

P = Those air toxics retained on filter material, either glass fiber or Teflon®, during sampling.

M = Airborne particulate with metallic constituents.

(4) Available standards.

+ U.S. EPA, Quality Assurance Division, AREAL, RTP, NC, Group 5/6 gas standards.

N Neat solution available from manufacturers.

G Gas cylinder standards produced and validated by consultants under EPA contract.

NT National Institute of Standards and Technology (NIST) solutions available.

F7 U.S. EPA, Quality Assurance Division, AREAL, RTP, NC, Future Group 7 gas standards.

(5) Notation

Sep-PAK® Silica gel impregnated with 2,4-Dinitrophenylhydrazine for extracting aldehydes and ketones from air.

Canister SUMMA® passivated stainless steel canister for collecting whole air samples.

Adsorbent Solid adsorbents, typically Tenax-GC

Filter Filter material, either glass fiber, Teflon or nylon, used to retain particles.

PUF Polyurethane foam for retaining semi-volatile pollutants.

IC Ion chromatography analysis using conductivity detector.

GC/MS Gas chromatography/mass spectroscopy analysis, applicable to both canisters and solid adsorbents.

ICAP Inductively coupled argon plasma spectroscopy analysis, applicable for metal analyses.

HPLC High performance liquid chromatography using ultraviolet detector.

(6) Available sampling/analytical notation

ACM Ambient continuous monitor.

ADS-AA Solid adsorbent sampling followed by flameless atomic adsorption analysis.

ADS-GC/MS Solid adsorbent sampling followed by gas chromatography/mass spectroscopy analysis.

C-C/MS Canister sampling followed by chromatography/mass spectroscopy analysis.

C-GC/MS Canister sampling by gas chromatography/mass spectroscopy analysis.

Abbreviations for Table G-1 (continued).

CT-GC/ECD	Activated charcoal tube sampling followed by gas chromatography with electron capture.
CT-GC/FID	Activated charcoal tube sampling followed by gas chromatography with flame ionization.
CT-GFAA	Activated charcoal tube adsorbent followed by radiochemistry
CV-AA	Filter sampling followed by cold vapor atomic adsorption spectroscopy.
DI-ICAP	Dichotomous sampling followed by inductively coupled argon plasma spectroscopy analysis.
Denuder-IC	Annual Denuder sampling followed by ion chromatographic analysis.
F-AA	Filter sampling followed by atomic adsorption spectroscopy.
F-GC/NPD	Filter sampling followed by gas chromatography separation with nitrogen-phosphorus detection.
F-GFAA	Filter sampling followed by graphite furnace atomic adsorption spectroscopy.
F-HPLC/UV	Filter sampling followed by high performance liquid chromatography with ultraviolet detection.
F-ICAP	Filter sampling followed by inductively coupled argon plasma spectroscopic analysis.
F-Micr	Filter sampling followed by microscopic analysis.
F/CT-GFAA	Filter/activated charcoal tube sampling with graphite furnace atomic absorption spectroscopy analysis.
F/CT-GC/FID	Filter/activated charcoal tube sampling followed by gas chromatography with flame ionization detection.
F/Imp-HPLC/UV	Filter/impinger sampling followed by high performance liquid chromatography with ultraviolet detection.
F/SG-GC/FID	Filter/silica gel adsorbent followed by gas chromatography with flame ionization detection.
F/SG-HPLC/UV	Filter/silica gel sorbent followed by high performance liquid chromatography with ultraviolet detection.
GB-GC/FID	Glass bulb sampling followed by gas chromatography separation with flame ionization detection.
GB-GC/FPD	Glass bulb sampling followed by gas chromatography separation with flame photometric detection.
GB-GC/MS	Glass bulb sampling followed by gas chromatography separation with mass spectroscopy identification.
Imp-COL	Impinger sampling followed by colorimetric analysis.
Imp-HPLC	Impinger sampling followed by high performance liquid chromatography.

PUF-GC/ECD	Polyurethane foam of XAD-2 sampling followed by a gas chromatography separation with electron capture detection.
PUF-GC/FID	Polyurethane foam sampling followed by gas chromatography separation with flame ionization detection.

Abbreviations for Table G-1 (continued).

PUF-GC/FPD	Polyurethane foam sampling followed by gas chromatography separation with flame photometric detection.
PUF-GC/MS	Polyurethane foam sampling followed by gas chromatography/mass spectroscopy analysis.
PUF/XAD-2-GC/MS	Polyurethane foam combined with XAD-2 resin for sampling followed by gas chromatography/mass spectroscopy analysis.
PUF-GC/NPD	Polyurethane foam sampling followed by high performance liquid chromatography.
PUF-HRGC/HRMS	Polyurethane foam sampling followed by high resolution gas chromatography with high resolution mass spectroscopy
S(Chromo 104)-GC/FID	Sorbent (chromosorb 104) sampling followed by gas chromatography separation with high resolution mass spectroscopy.
S(firbk)-HPLC/UV	Sorbent (firebrick) sampling followed by high performance liquid chromatography analysis.
S(Porapak-QS)-GC/NPD	Sorbent (Porapak-QS) sampling followed by gas chromatography separation with nitrogen-phosphorus detection.
S(silica gel)-GC/FID	Adsorbent (silica gel) sampling followed by gas chromatography separation with flame ionization detection.
S(silica gel)-GC/FID	Adsorbent (silica gel) sampling followed by gas chromatography separation with flame ionization detection.
S(silica gel)-HPLC/UV	Sorbent (silica gel) sampling followed by high performance liquid chromatography with ultraviolet detection.
S(TN)-GC/HECD	Sorbent (Thermosorb N) sampling followed by gas chromatography separation with Hall electron capture detector.
S(TN)-GC/NPD	Sorbent (Thermosorb N) sampling followed by gas chromatography separation with nitrogen phosphorus detection.
S(XAD-2)-HPLC/UV	Sorbent (XAD-2) sampling followed by high performance liquid chromatography analysis.
S(XAD-7)-HPLC/UV	Sorbent (XAD-7) sampling followed by high performance liquid chromatography analysis.
SEP-HPLC	Sep-PAK® impregnated cartridge sampling followed by high performance liquid chromatography.
SEP-IC	Sep-PAK® impregnated cartridge sampling followed by ion chromatography analysis.
T-GC/MS	Tenax solid adsorbent tube sampling followed by gas chromatography/mass spectroscopy analysis.

T/C-GC/MS Tenax solid adsorbent tube or canister sampling followed by gas chromatography/
mass spectroscopy analysis.

(7) Detection limits

- (a) Not amenable to Tenax analysis.
- (b) Not amenable to canister analysis.
- (c) No detection limits available, but feasible.
- (d) Canister - GC/MS in the SIM mode, Hewlett-Packard 5988A, column: 30 m x 0.32 i.d., DB-624 fused silica capillary, Perma Pure Dryer, 200 mL cryotrap sample, seven replicate samples analysis, LDD = (std. DEV.) x (one-tailed Student's value at 99% level).
- (e) Detection limit based upon 2500 m³ of air sampled, through a 8" x 10" glass filter with a 0.75" x 1" strip analyzed in final sample volume of 40 mL acid extraction solution.
- (f) PUF - Amount of air sampled determines MDLs. MDL based upon 273 m³ of theoretical air sampled, evaporate to 1 mL and analyze 1 µL by GC/MS/SIM.

(8) Approximate Air Risk Specific Concentration = [Acceptable Risk Level (i.e., 10⁻⁶)]/[Unit Risk Factor].

(9) Reference Ambient Levels (RALs) were developed from state agency acceptable ambient levels (AALs) as approximations of potential Applicable or Relevant and Appropriate Requirements (ARARs) or "To-Be-Considered" materials (TBCs) in establishment of air cleanup standards for remedial actions at national Priority List (NPL) sites.